

Competition of magnetism and superconductivity in underdoped $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$

To cite this article: Marianne Rotter *et al* 2009 *New J. Phys.* **11** 025014

View the [article online](#) for updates and enhancements.

Related content

- [Synthesis, crystal structure and SDW anomaly of the iron arsenide-fluoride \$\text{SrFeAsF}\$](#)
M. Tegel, S. Johansson, V. Weiß *et al.*
- [Local structure and hyperfine interactions of \$^{57}\text{Fe}\$ in \$\text{NaFeAs}\$ studied by Mössbauer spectroscopy](#)
I Presniakov, I Morozov, A Sobolev *et al.*
- [Structural and magnetic phase transitions in the ternary iron arsenides \$\text{SrFe}_2\text{As}_2\$ and \$\text{EuFe}_2\text{As}_2\$](#)
Marcus Tegel, Marianne Rotter, Veronika Weiß *et al.*

Recent citations

- [Evidence for negative thermal expansion in the superconducting precursor phase \$\text{SmFeAsO}\$](#)
H D Zhou *et al*
- [Specific Heat of \$\text{Ba}_{0.59}\text{K}_{0.41}\text{Fe}_2\text{As}_2\$, an Fe-Pnictide Superconductor with \$T_c = 36.9\$ K, and a New Method for Identifying the Electron Contribution](#)
Costel R. Rotundu *et al*
- [Effect of spin-density wave fluctuations on the specific heat jump in iron pnictides at the superconducting transition](#)
D. Kuzmanovski *et al*

Competition of magnetism and superconductivity in underdoped $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$

Marianne Rotter¹, Marcus Tegel¹, Inga Schellenberg²,
Falko M Schappacher², Rainer Pöttgen², Joachim Deisenhofer³,
Axel Günther³, Florian Schrettle³, Alois Loidl³
and Dirk Johrendt^{1,4}

¹ Department Chemie und Biochemie der Ludwig-Maximilians-Universität
München, Butenandtstr.-5-13 (Haus D), 81377 München, Germany

² Institut für Anorganische und Analytische Chemie, Universität Münster,
Corrensstrasse 30, D-48149 Münster, Germany

³ Experimentalphysik V, Center for Electronic Correlations and Magnetism,
Institute for Physics, Augsburg University, D-86135 Augsburg, Germany
E-mail: johrendt@lmu.de

New Journal of Physics **11** (2009) 025014 (18pp)

Received 14 December 2008

Published 27 February 2009

Online at <http://www.njp.org/>

doi:10.1088/1367-2630/11/2/025014

Abstract. Polycrystalline samples of underdoped $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ ($x \leq 0.4$) were synthesized and studied by x-ray powder diffraction, magnetic susceptibility, specific heat and ^{57}Fe -Mössbauer spectroscopy. The structural phase transition from tetragonal to orthorhombic lattice symmetry shifts towards lower temperatures, becomes less pronounced at $x = 0.1$ – 0.2 and is no longer present at $x = 0.3$. Bulk superconductivity is observed in all samples except $(\text{Ba}_{0.9}\text{K}_{0.1})\text{Fe}_2\text{As}_2$ by resistivity and magnetic susceptibility measurements. Specific heat data show a broad spin-density-wave (SDW) phase transition in $(\text{Ba}_{0.9}\text{K}_{0.1})\text{Fe}_2\text{As}_2$, which is hardly discernible in $(\text{Ba}_{0.8}\text{K}_{0.2})\text{Fe}_2\text{As}_2$. No SDW anomaly is found in the specific heat of optimally doped $(\text{Ba}_{0.6}\text{K}_{0.4})\text{Fe}_2\text{As}_2$, where C changes by 0.1 J K^{-1} at $T_c = 37.3 \text{ K}$. ^{57}Fe -Mössbauer spectra show full magnetic hyperfine field splitting, indicative of antiferromagnetic (AF) ordering at 4.2 K in samples with $x = 0$ – 0.2 , but zero magnetic hyperfine field in samples with $x = 0.3$. The spectra of $(\text{Ba}_{0.9}\text{K}_{0.1})\text{Fe}_2\text{As}_2$ and $(\text{Ba}_{0.8}\text{K}_{0.2})\text{Fe}_2\text{As}_2$ in the phase transition regions are temperature-dependent superpositions of magnetic and non-magnetic components, caused by inhomogeneous potassium distribution. Our results suggest the co-existence of AF ordering and

⁴ Author to whom any correspondence should be addressed.

superconductivity without mesoscopic phase separation in the underdoped region and show unambiguously homogeneous superconducting phases close to optimal doping. This is in contrast to recently reported results about single-crystal $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$.

Contents

1. Introduction	2
2. Experimental	3
2.1. Sample preparation	3
2.2. X-ray structure determination	4
2.3. Magnetic susceptibility and specific heat	4
2.4. Mößbauer spectroscopy	4
3. Results and discussion	4
3.1. Crystal structures and phase transitions	4
3.2. dc resistivity	5
3.3. Magnetic susceptibility	6
3.4. Specific heat	8
3.5. ^{57}Fe -Mößbauer spectroscopy	10
4. Conclusion	15
Acknowledgments	17
References	17

1. Introduction

The discovery of superconductivity (SC) in iron arsenides [1, 2] with transition temperatures (T_c) up to 55 K [3] has attracted enormous interest in the scientific community [4]. Besides the outstanding physical properties of this new class of superconducting materials, scientists found new hope that iron arsenides may help to finally solve the mystery of high- T_c SC. But prior to this long-term objective, many fundamental issues of the iron arsenides need to be clarified. Among them, the structural and magnetic phase diagrams with respect to doping, reflecting the interplay between SC and magnetism, are discussed controversially.

In both the LaFeAsO (1111) and BaFe_2As_2 (122) families, SC evolves from poor metallic parent compounds with quasi two-dimensional tetragonal crystal structures, which are subject to orthorhombic lattice distortions below certain temperatures (T_o). Static long-range antiferromagnetic (AF) ordering emerges with Néel temperatures (T_N) well below T_o in LaFeAsO [5], but very close to T_o in BaFe_2As_2 [6]. The structural and magnetic transitions of the parent compounds are strongly affected by doping of the FeAs layers either with electrons or holes, and SC appears at certain doping levels. For the underdoped phases in the transition zone, it has been reported that SC and AF ordering is either separated or co-existing. Also, the overlap of the orthorhombic distortion with the SC and AF areas in the phase diagrams is still not clear, neither in the 1111 nor in the 122 systems.

The first phase diagram of $\text{LaFeAsO}_{1-x}\text{F}_x$, constructed by μSR data, showed a sharp-cut vertical border between the SC and the orthorhombic AF phases at $x = 0.045$ [7]. But neutron diffraction experiments showed that although the magnetic ordering vanishes around $x \approx 0.04$,

the orthorhombic lattice still exists at least to $x = 0.05$, where SC has already emerged [8]. This is in line with the recently published neutron study of $\text{CeFeAsO}_{1-x}\text{F}_x$, where AF ordering disappears exactly before SC emerges, but the orthorhombic lattice persists extensively into the SC dome up to $x \approx 0.1$ [9]. Similar results have been reported for $\text{SmFeAsO}_{1-x}\text{F}_x$ from μSR experiments [10] and structural investigations using synchrotron radiation [11]. Thus at the moment all signs are that in the case of the 1111-family, static AF order is completely suppressed before SC emerges, but the orthorhombic lattice co-exists with SC and the temperature difference between T_0 and T_N increases with the doping level. This behavior of the 1111-superconductors is strongly reminiscent of the monolayer high- T_c cuprates. For instance in $\text{La}_{2-x}\text{Sr}_x\text{CuO}_4$, the AF order is well separated from the SC state, but the orthorhombic phase exists largely in the superconducting dome [12].

In the 122-family, co-existence of the orthorhombic structure with SC was first published for $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ up to $x \approx 0.2$ ($T_c \approx 26$ K) by x-ray powder diffraction [13]. The following neutron diffraction experiments also showed orthorhombic symmetry and long-range AF ordering co-existing up to $x = 0.3$ ($T_c < 15$ K) [14]. The different shapes of the superconducting domes $T_c(x)$ of $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ may be due to different synthesis conditions. However, the x values in [13] are determined from x-ray data by Rietveld refinements, whereas only the nominal compositions are given in [14]. Since diffraction methods provide the mean structural information on a rather long spatial scale, short-range phase inhomogeneities are averaged. Thus one may understand the observed co-existence of SC with AF ordering in $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ by phase separation in magnetic non-superconducting and non-magnetic superconducting mesoscopic domains. Local probes such as μSR and ^{57}Fe -Mössbauer spectroscopy can provide more accurate information. Recently, three reports about μSR experiments, each conducted with almost optimally doped superconducting $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ single crystals, concluded consistently phase separations into SC and AF domains. The non-magnetic superconducting volume fractions were found to be $\approx 30\%$ [15], 40% [16] and 25% [17]. In the latter report, the lateral scale of the inhomogeneities was estimated to 65 ± 10 nm by magnetic force microscopy (MFM) imaging. However, the onset of AF ordering in the superconducting crystals was detected at ≈ 70 – 80 K, irrespective of different doping levels.

In the present paper, we report on a detailed study of the structural and magnetic transitions of polycrystalline underdoped $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ ($x \leq 0.4$). The samples were characterized by magnetic susceptibility and specific heat measurements. The crystal structures and chemical compositions were determined by Rietveld refinements of x-ray powder patterns. Detailed temperature-dependent ^{57}Fe -Mössbauer spectra were recorded in order to detect the evolution of magnetic ordering on a local spatial scale.

2. Experimental

2.1. Sample preparation

Samples of $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ with $x = 0.1, 0.2$ and 0.3 were prepared by heating stoichiometric mixtures of the elements (all purities $> 99.9\%$) in alumina crucibles enclosed in silica tubes under an atmosphere of purified argon. In order to minimize the loss of potassium by evaporation at elevated temperatures, the gas volume in the crucibles was reduced by alumina inlays. The mixtures were heated slowly (50 K h^{-1}) to 873 K, kept at this temperature for 15 h and cooled down to room temperature. The reaction products were homogenized in an agate mortar and

annealed at 923 K for 15 h. After cooling, the samples were homogenized again, pressed into pellets and sintered at 1023 K for 15 h. The obtained black crystalline powders are stable in air for weeks. The Ba : K ratios were checked by EDX and chemical analysis (ICP-AAS), which resulted in the nominal composition within 5%.

2.2. X-ray structure determination

Phase purity was checked by x-ray powder diffraction using a Huber G670 Guinier imaging plate diffractometer (Cu- K_{α_1} radiation, Ge-111 monochromator), equipped with a closed-cycle He-cryostat. Rietveld refinements of all diffractograms were performed with the TOPAS package [18] using the fundamental parameters approach as reflection profiles (convolution of appropriate source emission profiles with equatorial and axial instrument contributions as well as crystallite microstructure effects). In order to obtain crystal structures inclusive of the Ba : K ratios, all profile contributions were refined freely, but in order to obtain accurate lattice parameter changes, all profile contributions were refined at room temperature and held constant for all other temperatures (except for the Lorentzian strain contribution). All diffractograms were measured without an internal standard, so the absolute lattice parameters might be slightly offset. In all cases, an empirical 2θ -dependent intensity correction for different absorption lengths arising from the Guinier geometry setup was applied.

2.3. Magnetic susceptibility and specific heat

The magnetic properties were studied using a commercial SQUID magnetometer (Quantum Design MPMS-5). The heat capacity was measured in a Quantum Design Physical Property Measurement System for temperatures from 1.8 to 300 K and magnetic fields up to 9 T.

2.4. Mößbauer spectroscopy

A $^{57}\text{Co}/\text{Rh}$ source was available for the ^{57}Fe -Mößbauer spectroscopy investigations. The samples were placed in thin-walled PVC containers at a thickness of about 10 mg Fe cm^{-2} . The measurements were run in the usual transmission geometry in the temperature range from room temperature to 4.2 K. The source was kept at room temperature. The total counting times per spectrum ranged between 5 h and 1 day.

3. Results and discussion

3.1. Crystal structures and phase transitions

The crystal structures were determined by x-ray powder diffraction. Figure 1 shows the x-ray powder patterns of $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ ($x = 0.1, 0.2, 0.3$) at 300 K with Rietveld-fits and the difference lines. Crystallographic data and selected bond lengths and angles at 300 and 10 K, respectively, are compiled in table 1. The temperature dependences of the a and b lattice parameters are shown in figure 2. In line with [13], the parameter a of the tetragonal phase decreases with the doping level x , while c increases (not shown). The tetragonal-to-orthorhombic phase transition is strongly affected by the potassium content. The transition temperature T_0 decreases to ≈ 100 K at $x = 0.2$ and is no longer visible at $x = 0.3$. Also the magnitude of the distortion, expressed by the differences between a and b at 10 K, decreases

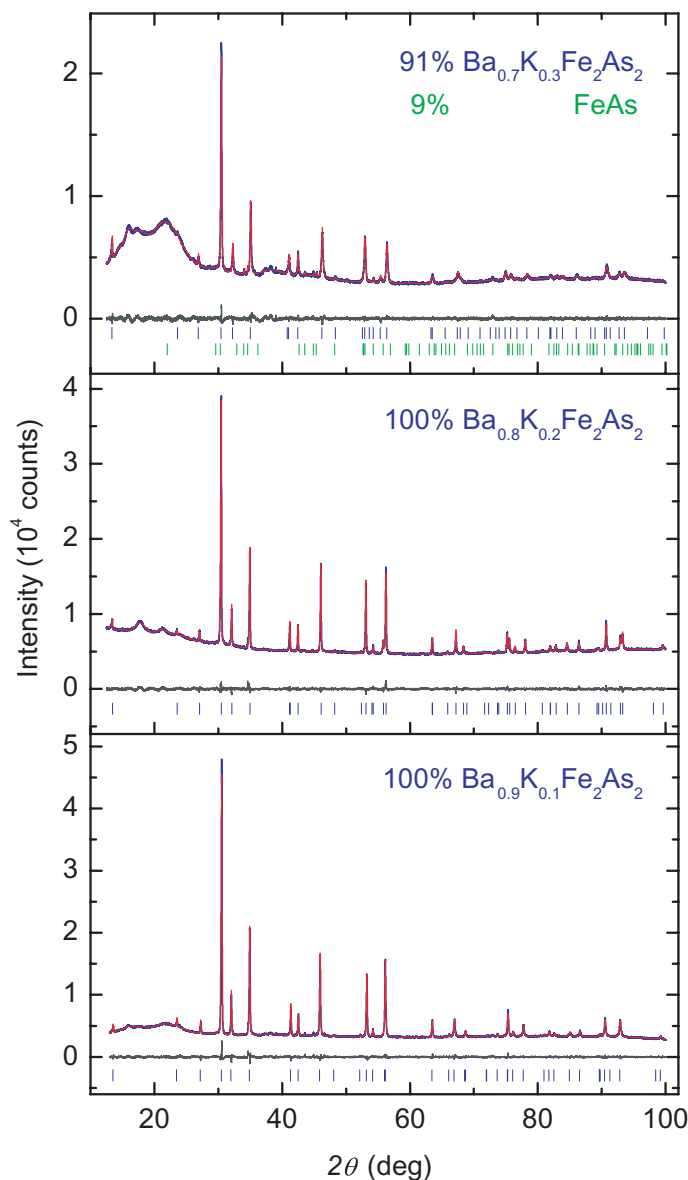


Figure 1. X-ray powder patterns of $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ ($x = 0.1, 0.2, 0.3$) with Rietveld profile fits and difference lines.

from 0.73% ($x = 0$) to 0.70% ($x = 0.1$) to 0.49% ($x = 0.2$). Thus with increasing potassium doping levels, the structural transition of BaFe_2As_2 is shifted towards lower temperatures and also less pronounced. It is no longer present at $x = 0.3$ (see figure 2).

3.2. *dc* resistivity

The temperature dependence of the *dc* resistivity of $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ is shown in figure 3. At the lowest doping concentration ($x = 0.1$), the typical SDW anomaly is still visible, but shifted towards lower temperatures and less pronounced than in undoped BaFe_2As_2 [6]. We observe a drop of the resistance below 3 K, associated with a superconducting transition, even though zero

Table 1. Crystal structure data of $\text{Ba}_{1-x}\text{K}_x\text{Fe}_2\text{As}_2$ at different temperatures.

	$x = 0.1$		$x = 0.2$		$x = 0.3$
Temp. (K)	300	10	300	10	300
Space group	$I4/mmm$	$Fmmm$	$I4/mmm$	$Fmmm$	$I4/mmm$
a (pm)	395.37(1)	560.07(1)	393.95(1)	557.34(1)	392.57(1)
b (pm)	$= a$	556.20(1)	$= a$	554.64(1)	$= a$
c (pm)	1310.60(1)	1301.35(1)	1318.90(3)	1309.26(3)	1327.02(3)
V (nm ³)	0.204 87(1)	0.405 38(1)	0.204 69(1)	0.404 72(1)	0.204 51(1)
Z	2	4	2	4	2
Data points	17 401	17 401	17 401	17 401	17 501
Reflections	46	70	46	70	46
d range	1.012–6.553	1.006–6.507	1.009–6.595	1.005–6.546	1.007–6.635
R_P, wR_P	0.0153, 0.0211	0.0154, 0.0210	0.0158, 0.0217	0.0172, 0.0229	0.0146, 0.0192
R_{Bragg}, χ^2	0.0121, 1.319	0.0132, 1.308	0.0123, 1.190	0.0117, 1.222	0.0076, 1.212
K, Ba	$2a$ (0,0,0) $U_{\text{iso}} = 119(4)$	$2a$ (0,0,0) $U_{\text{iso}} = 45(4)$	$2a$ (0,0,0) $U_{\text{iso}} = 173(7)$	$2a$ (0,0,0) $U_{\text{iso}} = 117(8)$	$2a$ (0,0,0) $U_{\text{iso}} = 181(6)$
Fe	$4d$ ($\frac{1}{2}, 0, \frac{1}{4}$) $U_{\text{iso}} = 114(4)$	$8f$ ($\frac{1}{4}, \frac{1}{4}, \frac{1}{4}$) $U_{\text{iso}} = 43(4)$	$4d$ ($\frac{1}{2}, 0, \frac{1}{4}$) $U_{\text{iso}} = 156(7)$	$8f$ ($\frac{1}{4}, \frac{1}{4}, \frac{1}{4}$) $U_{\text{iso}} = 102(8)$	$4d$ ($\frac{1}{2}, 0, \frac{1}{4}$) $U_{\text{iso}} = 58(5)$
As	$4e$ (0,0, z) $z = 0.3547(1)$ $U_{\text{iso}} = 147(4)$	$8i$ (0,0, z) $z = 0.3538(1)$ $U_{\text{iso}} = 57(4)$	$4e$ (0,0, z) $z = 0.3545(1)$ $U_{\text{iso}} = 129(7)$	$8i$ (0,0, z) $z = 0.3537(1)$ $U_{\text{iso}} = 72(8)$	$4e$ (0,0, z) $z = 0.3545(1)$ $U_{\text{iso}} = 79(5)$
K : Ba ratio	14(1) : 86(1)	13(1) : 87(1)	20(1) : 80(1)	20(1) : 80(1)	24(1) : 76(1)
Lengths (pm):					
Ba–As	338.3(1) \times 8	337.0(1) \times 4 338.6(1) \times 4	338.3(1) \times 8	337.0(1) \times 4 338.2(1) \times 4	337.7(1) \times 8
Fe–As	240.6(1) \times 4	239.1(1) \times 4	240.4(1) \times 4	238.9(1) \times 4	240.8(1) \times 4
Fe–Fe	279.6(1) \times 4	278.1(1) \times 2 280.0(1) \times 2	278.6(1) \times 4	277.3(1) \times 2 278.7(1) \times 2	277.6(1) \times 4
Angles (deg):					
As–Fe–As	110.5(1) \times 2 109.0(1) \times 4	111.2(1) \times 2 108.9(1) \times 2 108.3(1) \times 2	110.1(1) \times 2 109.2(1) \times 4	110.7(1) \times 2 109.0(1) \times 2 108.6(1) \times 2	109.2(1) \times 2 109.6(1) \times 4

resistance could not be reached at 1.8 K. The curvature of the resistivity of $(\text{Ba}_{0.8}\text{K}_{0.2})\text{Fe}_2\text{As}_2$ is still reminiscent of an SDW anomaly, but smeared over a larger temperature range between ≈ 120 and 70 K. The superconducting transition at 24 K is rather broad (≈ 4 K), but zero resistance is clearly observed at 23 K. At the higher doping level $x = 0.3$, the resistivity shows no indications of the SDW anomaly and superconductivity emerges at $T_c = 33$ K.

3.3. Magnetic susceptibility

Field-cooled (FC) and zero-field cooled (ZFC) cycles of the static magnetic volume susceptibility are shown as a function of temperature in a magnetic field of 5 Oe in figure 4. Estimating the superconducting volume fractions for $x = 0.2$ and 0.4 from the ZFC value at 1.8 K as 0.93 and 0.94, respectively, bulk SC is evidently present. The FC values at

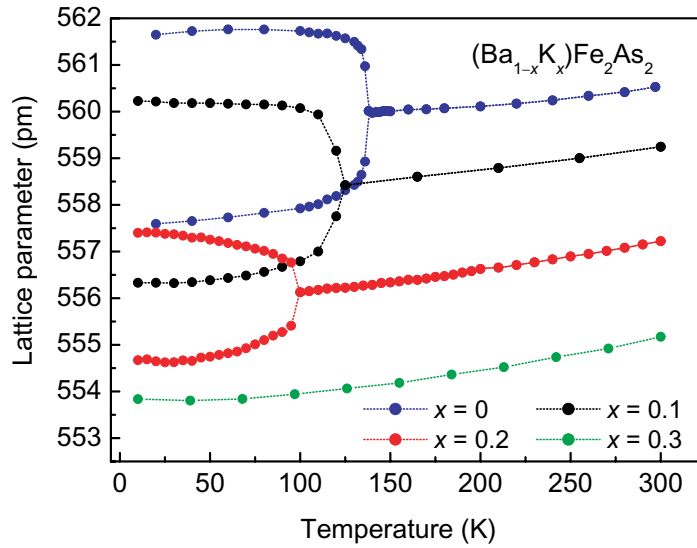


Figure 2. Lattice parameters of $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ ($x = 0, 0.1, 0.2, 0.3$). The tetragonal parameters are multiplied by $\sqrt{2}$ for comparison.

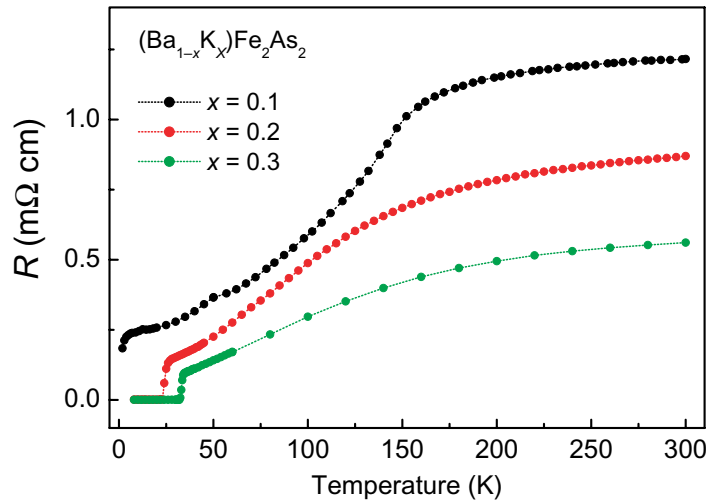


Figure 3. Temperature dependence of the dc resistivity of $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ ($x = 0.1, 0.2, 0.3$).

1.8 K amount to 1 and 64%, respectively. The corresponding temperatures, where 10% of the maximum shielding is reached, are $T_c = 23.6$ and 37.5 K for these two doping levels. For the sample with $x = 0.1$ bulk SC cannot be established, but the sample becomes diamagnetic below 5 K in the ZFC cycle as shown in figure 4(a). The constant positive contribution to the susceptibility may be due to ferromagnetic impurities. Therefore, we measured the complex ac-volume susceptibility $\chi_{ac} = \chi' + i\chi''$ for $x = 0.1$ with $H_{ac} = 0.5$ Oe and $f = 33$ Hz (see inset of figure 4(a)) and found that the sample exhibits a diamagnetic signal in χ' already below 11 K. Assuming that this diamagnetic signal is due to the onset of SC in the sample, this onset is strongly influenced by very weak external fields and points to a strong competition of SC and magnetism.

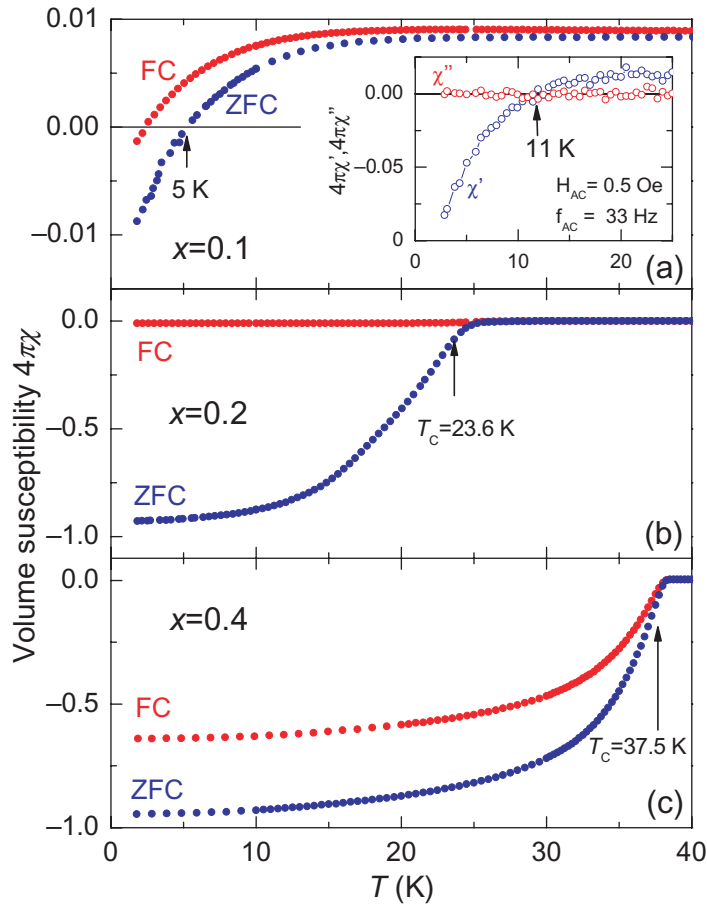


Figure 4. Static volume susceptibility $4\pi\chi = 4\pi M/H$ for $x = 0.1, 0.2$ and 0.4 in a magnetic field of 5 Oe for FC and ZFC cycles. Inset: real and imaginary parts of the ac volume susceptibility $4\pi\chi_{ac}$ measured at $H_{ac} = 0.5$ Oe and $f = 33$ Hz.

3.4. Specific heat

In figure 5 we show the temperature dependence of the specific heat for the optimally doped sample with $x = 0.4$. In the inset C_p/T is plotted as a function of temperature for zero magnetic field and a field of 9 T. Both curves are on top of each other above and below the superconducting transition. T_c in zero field is 37.3 K by the entropy conserving construction shown in the inset of figure 5. The change of the specific heat at the transition is estimated as $\Delta C|_{T_c} = 0.1 \text{ J mol K}^{-1}$, which is in good agreement with other reported values [19]–[21]. T_c is shifted by only 1 K in a field of 9 T reflecting the large upper critical fields, which were estimated as 70 T or even higher [20], [22]–[24].

The specific heat for the underdoped samples with $x = 0.1$ and 0.2 is plotted as a function of temperature in figure 6. For $x = 0.1$ one can clearly see a broad peak with a maximum at $T_0 = 132$ K, which is in agreement with the structural and magnetic transitions.

The low-temperature C/T data for the sample with $x = 0.1$ reveal the onset of a phase transition at $T_{\text{ons}} = 4.4$ K (inset of figure 6), which is completely suppressed in a magnetic field of 9 T. This temperature is in agreement with the appearance of diamagnetism as indicated in figure 4(a) and the appearance of a resistivity drop and therefore can be ascribed to the onset

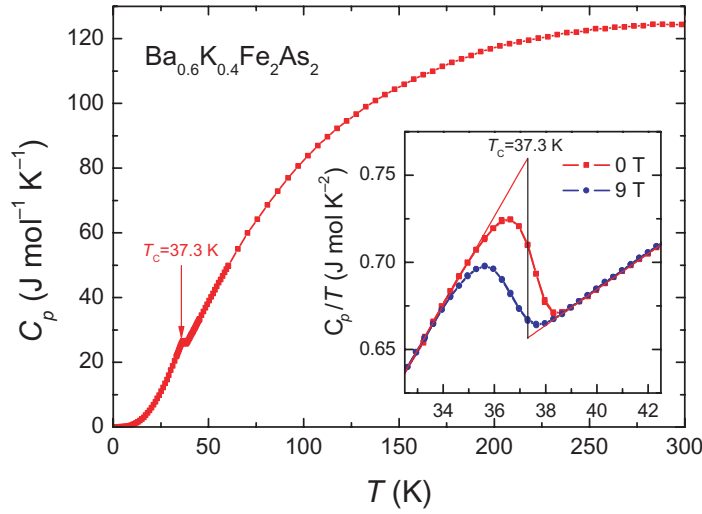


Figure 5. Specific heat of $\text{Ba}_{0.6}\text{K}_{0.4}\text{Fe}_2\text{As}_2$. Inset: comparison of C/T versus temperature in magnetic fields of 0 and 9 T. The lines are to guide the eyes.

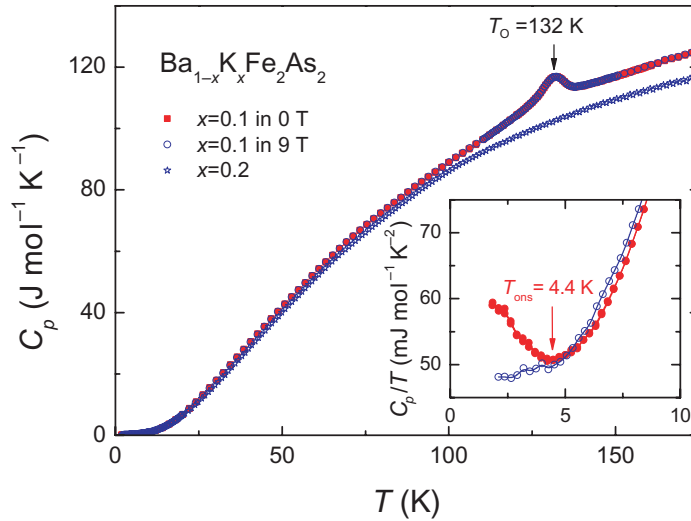


Figure 6. Comparison of C versus T for $x = 0.1$ and 0.2 . Inset: comparison of C/T versus T in magnetic fields of 0 and 9 T for $x = 0.1$ to reveal the onset of the superconducting transition and its suppression in a magnetic field of 9 T. The solid lines are to guide the eyes.

of SC. Hence, a superconducting and an AF transition are observable for $x = 0.1$, suggesting a strong competition of magnetic fluctuations and SC. However, one must keep in mind that bulk SC could not be established from susceptibility measurements and zero resistivity is not reached at 1.8 K.

Astonishingly, we could not detect any clearly visible anomaly for the sample with $x = 0.2$, where bulk SC is evident from figure 4(a) below $T_c = 23.6$ K and the structural transition occurs at $T_0 = 105$ K. In figure 7 we compare C/T versus T for these two concentrations and indicate the known transition temperatures for both compounds. While at around $T_0 = 105$ K,

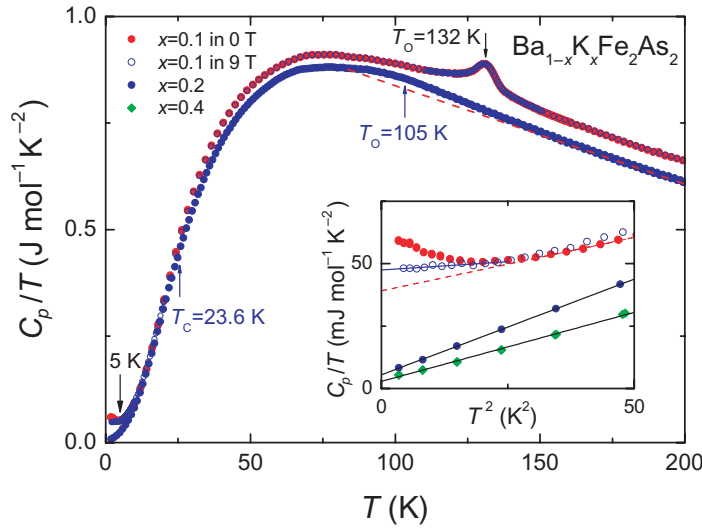


Figure 7. Comparison of C/T versus T for $x = 0.1$ and 0.2 . The dashed line is a guide for the eye. Inset: Comparison of C/T versus T^2 for $x = 0.1$ (0 and 9 T) and $x = 0.2, 0.4$ at lowest temperatures. The solid and dashed lines are linear fits described in the text.

an extremely broadened transition region between about 80 and 140 K may be visualized by guiding the eye with a linear extrapolation of the high-temperature behavior, a corresponding anomaly at the superconducting transition is not detectable in our data. The low-temperature data for all three concentrations $x = 0.1$ (in 0 and 9 T) and $x = 0.2, 0.4$ are plotted as C/T versus T^2 . We fitted the data for $x = 0.1$ at 9 T and for $x = 0.2, 0.4$ in zero magnetic field with linear behavior and extracted Sommerfeld coefficients of $\gamma = 47, 5.6, 2.8 \text{ mJ mol K}^{-2}$ and Debye temperatures $\Theta_D = 418, 238, 260 \text{ K}$, respectively (solid lines in the inset of figure 7). The values for γ and Θ_D for $x = 0.2$ and 0.4 are very close to each other and significantly lower than the values for $x = 0.1$. This reflects the superconducting low-temperature state for $x = 0.2$ and 0.4 , which is fully suppressed for $x = 0.1$ in a magnetic field of 9 T. Note that the value $\gamma = 47 \text{ mJ mol K}^{-2}$ and $\Theta_D = 418 \text{ K}$ may contain a magnetic-field-dependent contribution, because a linear fit (dashed line in the inset of figure 7) of the data above 5 K in zero field yield somewhat lower values of $\gamma = 39 \text{ mJ mol K}^{-2}$ and $\Theta_D = 283 \text{ K}$.

3.5. ^{57}Fe -Mössbauer spectroscopy

^{57}Fe -Mössbauer spectroscopy is an excellent local probe for the determination of magnetic ordering in iron compounds. In comparison with other experimental methods, Mössbauer spectroscopy studies of the new superconductors are rather rare. So far LaFePO [25], LaFeAsO [26]–[29], SrFeAsF [30], BaFe_2As_2 [6], SrFe_2As_2 [31] and EuFe_2As_2 [32] have been investigated. Since the formation of binary iron phosphide or arsenide impurities can seriously affect the property measurements, ^{57}Fe -Mössbauer spectroscopy can also be a useful analytical tool to detect iron impurity phases. The influence of Fe_2As , FeAs and FeAs_2 on the SDW transitions and the superconducting properties has been studied systematically [33]. First we discuss the spectra of BaFe_2As_2 and KFe_2As_2 . In our earlier report on the SDW anomaly of BaFe_2As_2 , ^{57}Fe -Mössbauer spectra had only been recorded at 298, 77 and 4.2 K [6]. We

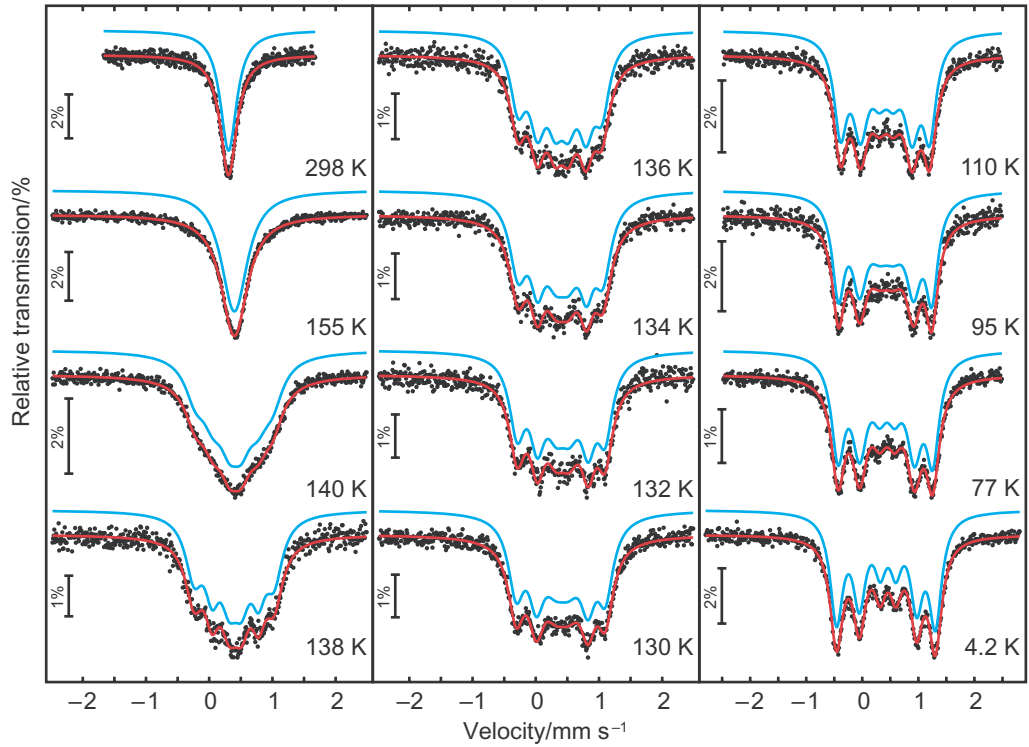


Figure 8. ^{57}Fe -Mössbauer spectra of BaFe_2As_2 .

have now studied BaFe_2As_2 over the entire temperature range. The spectra are presented in figure 8 together with transmission integral fits. The corresponding fitting parameters are listed in table 2. At room temperature, the spectrum consists of a single Lorentzian line with an isomer shift of $\delta = 0.31 \text{ mm s}^{-1}$. First hints at line broadening and thus short-range magnetic ordering appear already at 155 K, thus well above the structural distortion, however, with a very small magnetic hyperfine field (table 2). When lowering the temperature below the SDW transition temperature of 138 K, we observe a strong increase of the internal hyperfine field with a saturation value of 5.47 T at 4.2 K (see figure 8). This corresponds to a magnetic moment of approximately $0.4\text{--}0.5 \mu_{\text{B}}$ per iron atom. Similar behavior has been observed for SrFe_2As_2 [31] and the quaternary iron arsenide LaFeAsO [26]–[29]. In the latter two compounds, the saturation hyperfine fields are 8.9 T (SrFe_2As_2) and 5.3 T (LaFeAsO).

For BaFe_2As_2 reported herein and also for SrFe_2As_2 [31], the magnetically split spectra can be well reproduced by one spectral component with the typical sextet of lines. This is in contrast to the ZrCuSiAs -type compounds LaFeAsO [26] and SrFeAsF [30], both of which show distributions of moments in the magnetically ordered states, indicating that some spin disorder still remains. Most likely the ordering patterns in the ZrCuSiAs and ThCr_2Si_2 -type materials are slightly different.

The ^{57}Fe -Mössbauer spectra of KFe_2As_2 are shown in figure 9. KFe_2As_2 is a metal with a superconducting transition at $T_c = 3.8 \text{ K}$. We observe a single absorption line with an isomer shift of 0.21 mm s^{-1} from room temperature down to 4.2 K, which is subjected to weak quadrupole splitting (table 2). Since potassium transfers only one electron to the $[\text{Fe}_2\text{As}_2]$ layer, we expect a smaller electron density at the iron nuclei in KFe_2As_2 , which is consistent with

Table 2. Fitting parameters of the ^{57}Fe -Mössbauer spectroscopic measurements of BaFe_2As_2 , $\text{Ba}_{0.9}\text{K}_{0.1}\text{Fe}_2\text{As}_2$, $\text{Ba}_{0.8}\text{K}_{0.2}\text{Fe}_2\text{As}_2$ and KFe_2As_2 at different temperatures.

BaFe_2As_2

T (K)	δ_1 (mm s $^{-1}$)	ΔE_{Q1} (mm s $^{-1}$)	Γ_1 (mm s $^{-1}$)	B_{hf} (mm s $^{-1}$)
298	0.31(1)	0.00(1)	0.32(1)	—
155	0.40(1)	−0.06(22)	0.46(1)	0.37(58)
145	0.41(1)	−0.02(1)	0.30	1.79(9)
140	0.40(1)	−0.02(1)	0.39(4)	3.80(7)
138	0.40(1)	−0.02(1)	0.33(3)	3.93(4)
136	0.40(1)	−0.03(1)	0.32(2)	4.12(3)
134	0.40(1)	−0.02(1)	0.40(4)	4.16(3)
132	0.41(1)	−0.03(1)	0.38(3)	4.31(3)
130	0.41(1)	−0.02(1)	0.46(6)	4.37(2)
125	0.41(1)	−0.03(1)	0.48(6)	4.62(2)
110	0.42(1)	−0.02(1)	0.47(7)	5.16(1)
77	0.43(1)	−0.03(1)	0.33(2)	5.23(1)
4.2	0.44(1)	−0.04(1)	0.25(1)	5.47(1)

$(\text{Ba}_{0.9}\text{K}_{0.1})\text{Fe}_2\text{As}_2$

T (K)	δ_1 (mm s $^{-1}$)	ΔE_{Q1} (mm s $^{-1}$)	Γ_1 (mm s $^{-1}$)	B_{hf} (mm s $^{-1}$)	δ_2 (T)	ΔE_{Q2} (mm s $^{-1}$)	Γ_2 (mm s $^{-1}$)	A1/A2 (mm s $^{-1}$)
298					0.30(1)	−0.06(1)	0.28(1)	
160					0.40(1)	−0.04(2)	0.32(1)	
148	0.38(1)	−0.06(2)	0.32	3.32(8)	0.40(1)	—	0.30	28 : 72
146	0.41(1)	−0.02(1)	0.32	3.46(3)	0.39(1)	—	0.30	34 : 66
144	0.39(1)	−0.02(1)	0.32(1)	3.52(5)	0.39(1)	—	0.30	46 : 54
142	0.40(1)	−0.03(1)	0.33(1)	3.64(2)	0.40(1)	—	0.30	66 : 34
141	0.39(1)	−0.01(1)	0.30(1)	3.61(2)	0.39(1)	—	0.30	71 : 29
138	0.39(1)	−0.03(1)	0.39(1)	3.78(2)	0.38(1)	—	0.30	91 : 9
136	0.40(1)	−0.02(1)	0.37(1)	3.91(3)	0.38(1)	—	0.30	96 : 4
124	0.40(1)	−0.03(1)	0.29(1)	4.58(1)				
120	0.40(1)	−0.03(1)	0.28(1)	4.65(1)				
118	0.41(1)	−0.03(1)	0.29(1)	4.72(1)				
116	0.41(1)	−0.03(1)	0.28(1)	4.77(1)				
114	0.41(1)	−0.03(1)	0.28(1)	4.79(1)				
112	0.41(1)	−0.02(1)	0.28(1)	4.81(1)				
110	0.41(1)	−0.02(1)	0.27(1)	4.86(1)				
77	0.42(1)	−0.03(1)	0.27(1)	5.22(1)				
50	0.44(1)	−0.03(1)	0.31(1)	5.46(2)				
30	0.45(1)	−0.04(1)	0.36(1)	5.55(2)				
4.2	0.44(1)	−0.04(1)	0.36(1)	5.57(2)				

(Ba_{0.8}K_{0.2})Fe₂As₂

<i>T</i> (K)	δ_1 (mm s ⁻¹)	ΔE_{Q1} (mm s ⁻¹)	Γ_1 (mm s ⁻¹)	B_{hf} (mm s ⁻¹)	δ_2 (T)	ΔE_{Q2} (mm s ⁻¹)	Γ_2 (mm s ⁻¹)	A1/A2 (mm s ⁻¹)
298					0.30(1)	-0.09(1)	0.28(1)	
130					0.39(1)	0.00(1)	0.34(1)	
110	0.42(1)	-0.02(1)	0.26(2)	3.54(7)	0.39(1)	—	0.33	37 : 63
108	0.40(1)	-0.03(1)	0.28(2)	3.54(6)	0.41(1)	—	0.33	45 : 55
106	0.40(1)	-0.03(1)	0.29(2)	3.61(6)	0.41(1)	—	0.33	52 : 48
104	0.40(1)	-0.02(1)	0.30(2)	3.46(5)	0.40(1)	—	0.33	61 : 39
102	0.40(1)	-0.02(1)	0.30(2)	3.46(4)	0.40(1)	—	0.33	72 : 28
100	0.39(1)	-0.02(1)	0.29(1)	3.57(3)	0.40(1)	—	0.33	79 : 21
98	0.40(1)	-0.02(1)	0.29(1)	3.61(1)	0.39(1)	—	0.33	90 : 10
96	0.40(1)	-0.02(1)	0.28(1)	3.65(2)	0.37(1)	—	0.33	96 : 4
94	0.40(1)	-0.02(1)	0.27(3)	3.72(4)				
77	0.41(1)	-0.02(1)	0.29(3)	4.25(2)				
50	0.43(1)	-0.03(1)	0.33(4)	4.85(1)				
30	0.43(1)	-0.04(1)	0.30	5.07(3)				
4.2	0.43(1)	-0.04(1)	0.31(3)	5.07(2)				

KFe₂As₂

<i>T</i> (K)	δ_1 (mm s ⁻¹)	ΔE_{Q1} (mm s ⁻¹)	Γ_1 (mm s ⁻¹)
298	0.21(1)	-0.03(11)	0.39(1)
77	0.33(1)	-0.08(1)	0.33
4.2	0.34(1)	-0.09(4)	0.40(2)

Numbers in parentheses are the statistical errors in the last digit. Values without standard deviations were kept fixed during the fitting procedure. (δ), isomer shift; (Γ), experimental line width; (ΔE_Q), quadrupole splitting parameter; (B_{hf}), magnetic hyperfine field. A1/A2 is the ratio of the signals.

lower isomer shift compared to BaFe₂As₂ [6] and SrFe₂As₂ [31]. From this it is evident that the potassium doping affects the electronic situation of the iron atoms.

The potassium-doped samples (Ba_{1-x}K_x)Fe₂As₂ with $x = 0.1$ and 0.2 show strikingly different temperature dependences of the ⁵⁷Fe-Mössbauer spectra, shown in figures 10 and 11. In the temperature ranges 148–136 K ($x = 0.1$) and 110–96 K ($x = 0.2$), the spectra show superpositions of one magnetically split and one unsplit component. The non-magnetic components rapidly diminish within small temperature ranges (≈ 10 K). Below 136 and 96 K, respectively, the spectra of (Ba_{0.9}K_{0.1})Fe₂As₂ and (Ba_{0.8}K_{0.2})Fe₂As₂ can be well reproduced by one single magnetically split signal. These magnetic transition temperatures are very close to the structural transition temperatures extracted from the x-ray data. Similar to pure BaFe₂As₂, the magnetic hyperfine fields increase with decreasing temperature. BaFe₂As₂ and (Ba_{0.9}K_{0.1})Fe₂As₂ show almost similar hyperfine fields of ≈ 5.5 T at 4.2 K (table 2), while a decrease of B_{hf} by 10% is observed for (Ba_{0.8}K_{0.2})Fe₂As₂.

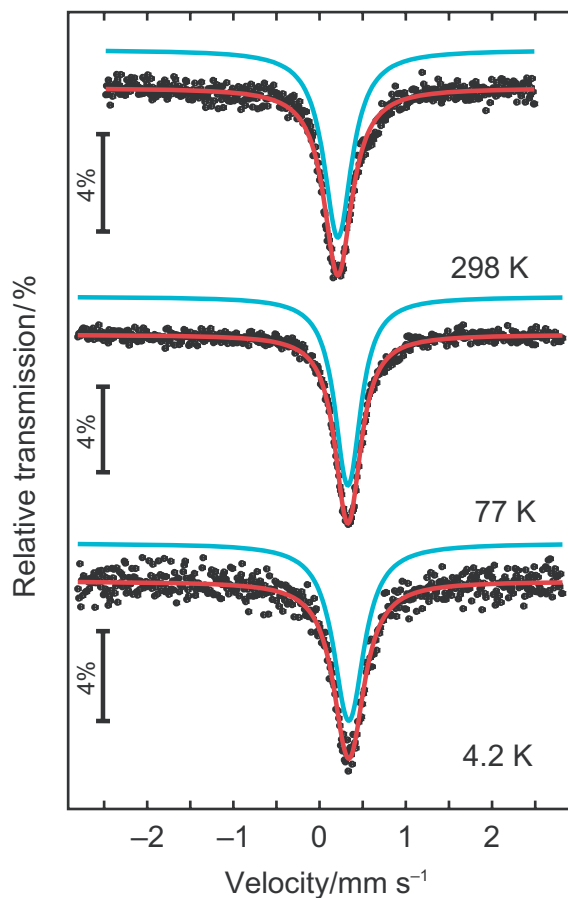


Figure 9. ^{57}Fe -Mössbauer spectra of KFe_2As_2 .

Table 3. Fitting parameters of the ^{57}Fe -Mössbauer spectroscopic measurements of $(\text{Ba}_{0.8}\text{K}_{0.2})\text{Fe}_2\text{As}_2$ and $(\text{Ba}_{0.7}\text{K}_{0.3})\text{Fe}_2\text{As}_2$ at 4.2 K.

	T (K)	δ_1 (mm s $^{-1}$)	ΔE_{Q1} (mm s $^{-1}$)	Γ_1 (mm s $^{-1}$)
$(\text{Ba}_{0.7}\text{K}_{0.3})\text{Fe}_2\text{As}_2$	4.2	0.41(1)	−0.02*	0.47(1)
$(\text{Ba}_{0.6}\text{K}_{0.4})\text{Fe}_2\text{As}_2$	4.2	0.39(1)	−0.10(1)	0.35(1)

These findings suggest that the spectra of the underdoped compounds ($x = 0.1$ and 0.2) are caused by temperature-dependent superpositions of paramagnetic and antiferromagnetically ordered domains. This reflects a chemical inhomogeneity of the Ba/K distribution, where the unsplit components represent domains with higher potassium content and lower Néel temperatures and vice versa. This is in agreement with the smaller isomer shifts of the unsplit signals, which indicate higher doping. On cooling, more domains get magnetically ordered until the paramagnetic fractions are completely consumed. Thus we see no distinct phase separation, but a continuous (narrow) distribution of the potassium concentration. Since no paramagnetic component exists below ≈ 136 and ≈ 96 K, respectively, we observe homogenous co-existence of AF magnetic ordering with superconductivity in $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ at $x = 0.1$ and 0.2 .

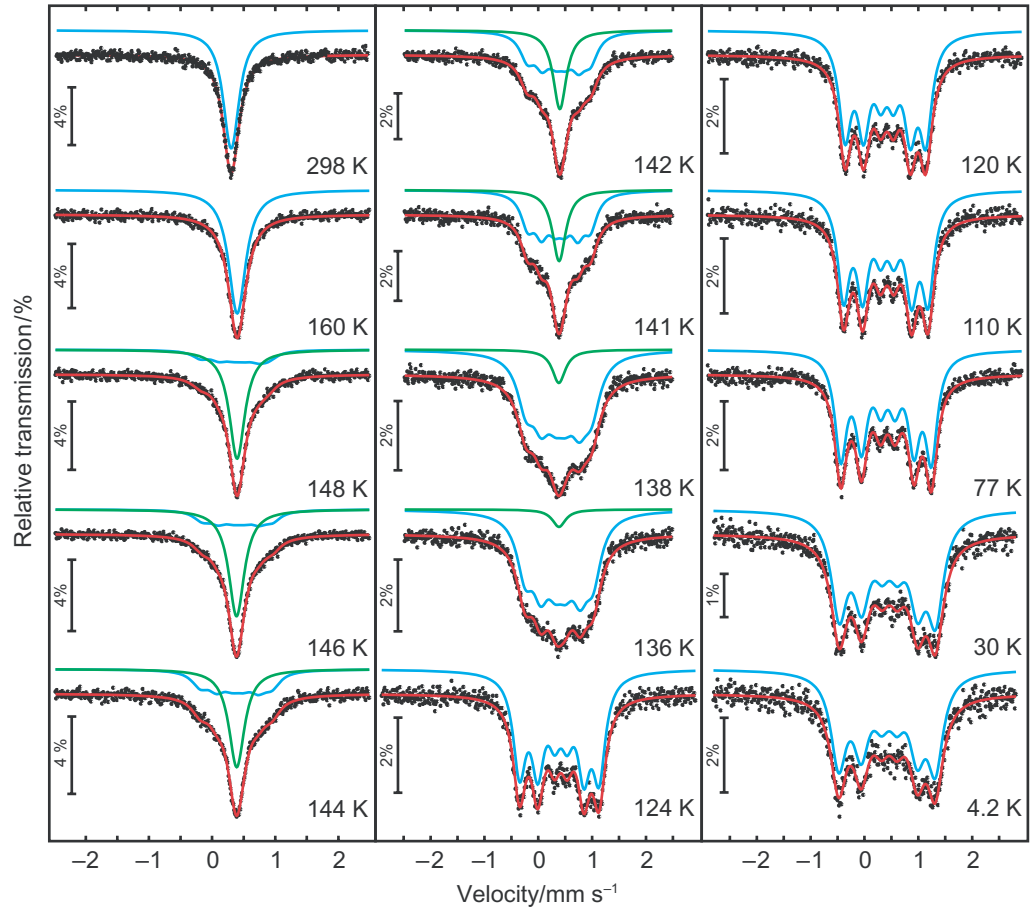


Figure 10. ^{57}Fe -Mössbauer spectra of $(\text{Ba}_{0.9}\text{K}_{0.1})\text{Fe}_2\text{As}_2$.

A further increase of doping again radically changes the ^{57}Fe -Mössbauer spectra. Figure 12 shows the spectra of $(\text{Ba}_{0.7}\text{K}_{0.3})\text{Fe}_2\text{As}_2$ ($T_c = 33$ K) and $(\text{Ba}_{0.6}\text{K}_{0.4})\text{Fe}_2\text{As}_2$ ($T_c = 38$ K) recorded at 4.2 K together with transmission integral fits. The fitting parameters are listed in table 3. In both cases we observe only unsplit absorption lines, which can be fitted by one component without any magnetic hyperfine field. The line width is slightly increased in comparison with undoped BaFe_2As_2 , hinting at a small inhomogeneity of the Ba/K concentrations. Thus, in contrast to the completely magnetically ordered phase detected at $x = 0.2$, we find the phases at $x = 0.3$ and 0.4 to be completely non-magnetic at low temperatures. This is in line with the absence of lattice distortions, since both compounds keep the tetragonal structure at low temperatures (see figure 2 and [13]).

4. Conclusion

In summary, we have studied the doping dependences of the physical properties of $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ in the underdoped region. The SDW anomaly connected with a structural phase transition is continuously suppressed by increasing doping concentrations and no longer observed in $(\text{Ba}_{0.7}\text{K}_{0.3})\text{Fe}_2\text{As}_2$. Specific heat measurements reveal a smearing of the

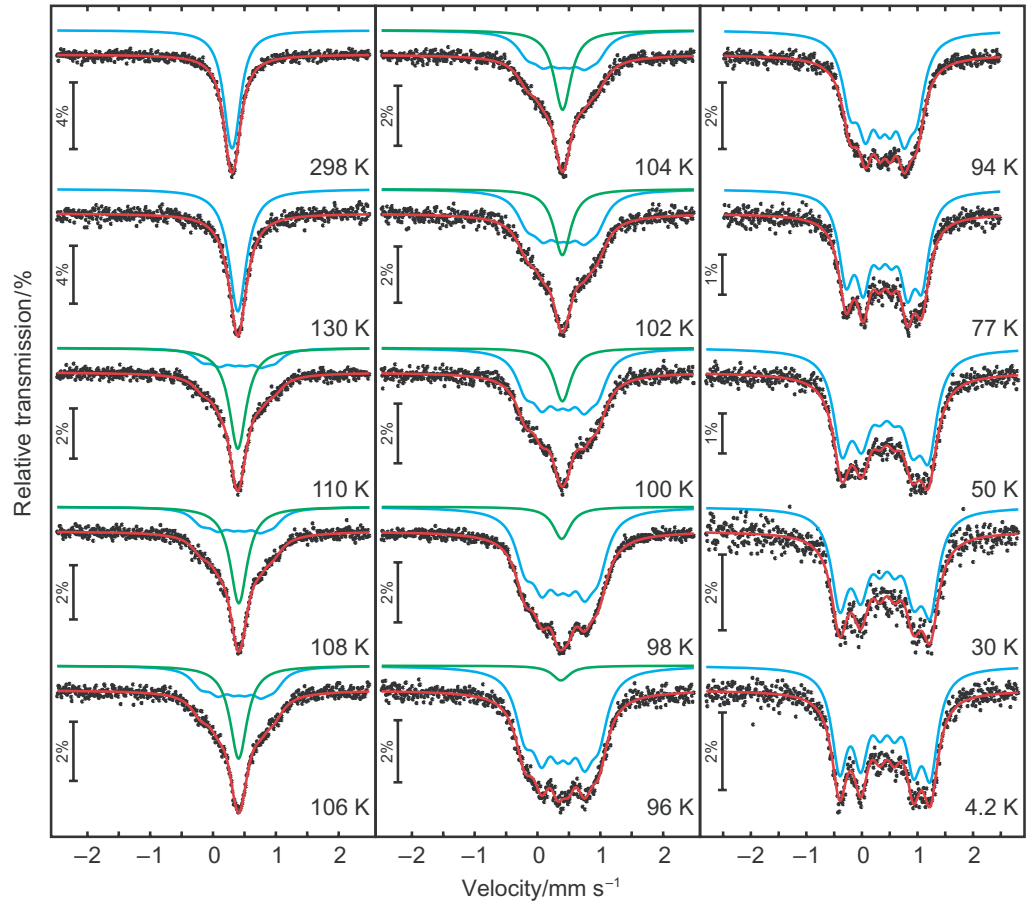


Figure 11. ^{57}Fe -Mössbauer spectra of $(\text{Ba}_{0.8}\text{K}_{0.2})\text{Fe}_2\text{As}_2$.

phase transitions over larger temperature ranges in the underdoped samples, but no SDW anomaly in the optimally doped compounds. This is in agreement with the structural data. Bulk SC was detected in all samples except $(\text{Ba}_{0.9}\text{K}_{0.1})\text{Fe}_2\text{As}_2$. ^{57}Fe -Mössbauer spectra of the underdoped compounds $(\text{Ba}_{0.9}\text{K}_{0.1})\text{Fe}_2\text{As}_2$ and $(\text{Ba}_{0.8}\text{K}_{0.2})\text{Fe}_2\text{As}_2$ show temperature-dependent superpositions of paramagnetic and antiferromagnetically ordered domains, which reflect chemical inhomogeneities of the Ba/K distributions resulting in different Néel temperatures of the domains. At lower temperatures, more domains get magnetically ordered until the paramagnetic fractions are completely consumed. Thus we see no distinct phase separation, but a continuous distribution of the potassium concentration. No paramagnetic component is observed in $(\text{Ba}_{0.8}\text{K}_{0.2})\text{Fe}_2\text{As}_2$ below T_c (24 K), suggestive of the co-existence of SC and AF ordering. Only at higher doping levels ($x = 0.3$) are the magnetic and structural phase transitions completely suppressed and SC reaches the highest T_c .

Our results contradict recent reports on mesoscopic phase separations in AF ordered and non-magnetic SC regions in single crystals of almost optimally doped $(\text{Ba}_{1-x}\text{K}_x)\text{Fe}_2\text{As}_2$ [15]–[17]. In the polycrystalline material studied here, the structural distortion and AF ordering are definitely absent already at $x = 0.3$. The origin of magnetically ordered fractions detected in almost optimally doped single crystals of $(\text{Ba}_{0.6}\text{K}_{0.4})\text{Fe}_2\text{As}_2$ with high T_c may either be attributed to a strongly inhomogeneous potassium distribution caused by

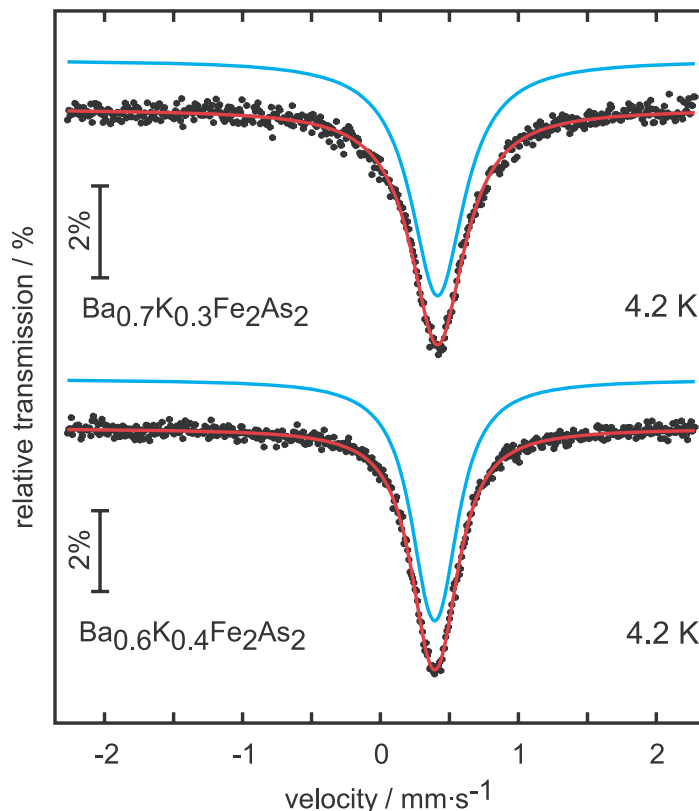


Figure 12. ^{57}Fe -Mössbauer spectra of $(\text{Ba}_{0.7}\text{K}_{0.3})\text{Fe}_2\text{As}_2$ and $(\text{Ba}_{0.6}\text{K}_{0.4})\text{Fe}_2\text{As}_2$ at 4.2 K.

uncontrolled single crystal growth or by magnetic impurity phases such as FeAs with a Néel temperature of 77 K [34].

Acknowledgments

It is a pleasure to thank E-W Scheidt and C Kant for fruitful discussions and A Pimenova for technical support. We acknowledge support by the BMBF via contract number VDI/EKM 13N6917 and by the DFG via SFB 484 (Augsburg) and project Jo257/5-1 (München).

References

- [1] Kamihara Y, Watanabe T, Hirano M and Hosono H 2008 *J. Am. Chem. Soc.* **130** 3296–7
- [2] Rotter M, Tegel M and Johrendt D 2008 *Phys. Rev. Lett.* **101** 107006
- [3] Ren Z *et al* 2008 *Chin. Phys. Lett.* **25** 2215
- [4] Johrendt D and Pöttgen R 2008 *Angew. Chem., Int. Edn Engl.* **47** 4782
- [5] de la Cruz C *et al* 2008 *Nature* **453** 899
- [6] Rotter M, Tegel M, Johrendt D, Schellenberg I, Hermes W and Pöttgen R 2008 *Phys. Rev. B* **78** 020503
- [7] Luetkens H *et al* 2008 arXiv:0806.3533
- [8] Huang Q, Qiu Y, Bao W, Lynn J, Green M, Chen Y, Wu T, Wu G and Chen X 2008 *Phys. Rev. B* **78** 054529
- [9] Zhao J *et al* 2008 *Nat. Mater.* **7** 953

- [10] Drew A J *et al* 2008 arXiv:0807.4876
- [11] Margadonna S, Takabayashi Y, McDonald M T, Brunelli M, Wu G, Liu R, Chen X H and Prassides K 2009 *Phys. Rev. B* **79** 014503
- [12] Keimer B *et al* 1992 *Phys. Rev. B* **46** 14034
- [13] Rotter M, Pangerl M, Tegel M and Johrendt D 2008 *Angew. Chem., Int. Edn Engl.* **47** 7949
- [14] Chen H *et al* 2009 *Europhys. Lett.* **85** 17006
- [15] Aczel A A *et al* 2008 arXiv:0807.1044
- [16] Goko T *et al* 2008 arXiv:0808.1425
- [17] Park J T *et al* 2008 arXiv:0811.2224
- [18] Cheary R W, Coelho A A and Cline J P 2004 *J. Res. Natl Inst. Stand. Technol.* **109** 1
- [19] Ni N *et al* 2008 *Phys. Rev. B* **78** 014507
- [20] Welp U, Xie R, Koshchev A E, Kwok W K, Luo H Q, Wang Z S, Mu G and Wen H H 2008 arXiv:0810.1944
- [21] Mu G, Luo H, Wang Z, Ren Z, Shan L, Ren C and Wen H-H 2008 arXiv:0812.1188
- [22] Wang Z S, Luo H-Q, Ren C and Wen H-H 2008 *Phys. Rev. B* **78** 140501
- [23] Yuan H Q, Singleton J, Balakirev F F, Baily S A, Chen G F, Luo J L and Wang N L X 2009 *Nature* **457** 565
- [24] Altarawneh M, Collar K, Mielke C H, Ni N, Bud'ko S L and Canfield P C 2008 *Phys. Rev. B* **78** 220505
- [25] Tegel M, Schellenberg I, Pöttgen R and Johrendt D 2008 *Z. Naturforsch. B* **63** 1057
- [26] Kitao S, Kobayashi Y, Higashitanguchi S, Saito M, Kamihara Y, Hirano M, Mitsui T, Hosono H and Seto M *J. Phys. Soc. Japan*
- [27] Klauss H H *et al* 2008 *Phys. Rev. Lett.* **101** 077005
- [28] Nowik I, Felner I, Awana V P S, Vajpayee A and Kishan H 2008 *J. Phys.: Condens. Matter* **20** 292201
- [29] McGuire M A *et al* 2008 *Phys. Rev. B* **78** 094517
- [30] Tegel M, Johansson S, Weiss V, Schellenberg I, Hermes W, Pöttgen R and Johrendt D 2008 *Europhys. Lett.* **84** 67007
- [31] Tegel M, Rotter M, Weiss V, Schappacher F M, Pöttgen R and Johrendt D 2008 *J. Phys.: Condens. Matter* **20** 452201
- [32] Raffius H, Mörsen E, Mosel B D, Müller-Warmuth W, Jeitschko W, Terbüchte L and Vomhof T 1993 *J. Phys. Chem. Solids* **54** 135
- [33] Nowik I and Felner I 2008 *J. Supercond. Nov. Magn.* **21** 297
- [34] Selte K, Kjekshus A and Andresen A F 1972 *Acta Chem. Scand.* **26** 3101